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File 410: The Chronolog 1991-2010/ Mar
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  File 5:Biosis Previews(R) 1926-2010/Jun W4
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  File 73:EMBASE 1974-2010/Jul 01
        (c) 2010 Elsevier B.V.
*File 73: The archive of Medline derived records was added to Embase.
  File 155:MEDLINE(R) 1950-2010/Jun 29
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*File 155: Medline has been reloaded. Please see HELP NEWS154
for information.
  File 399:CA SEARCH(R) 1967-2010/UD=15301
         (c) 2010 American Chemical Society
*File 399: Use is subject to the terms of your user/customer agreement.
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E4
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E5
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Ε6
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E7
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Ε2
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E5
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E9
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E10
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E12
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ts(w)a12?)
             111 S1
             135 S2
        2724863 ANTIBOD?
        1064219 IMMUNOGLOBULIN?
          61953 HYBRIDOMA?
          14523 C5A
          50236 TS
           5903 A12?
              4 TS(W)A12?
           1785 ((ANTIBOD? OR IMMUNOGLOBULIN?) OR HYBRIDOMA?) (20N) (C5A OR
                 TS(W)A12?)
      S3
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     S4
              2 RD S3 (unique items)
? t s4/3/all
          (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2010 The Thomson Corporation. All rts. reserv.
         BIOSIS NO.: 200200576587
16983076
The cleavage site of C5 from man and animals as a common target for
 neutralizing human monoclonal antibodies: In vitro and in vivo studies
AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti
 Fabio; Gennaro Renato; Marks James D; Bradbury Andrew; Tedesco
 Francesco (Reprint)
AUTHOR ADDRESS: Dipartimento di Fisiologia e Patologia, Universita di
  Trieste, Via Fleming 22, I-34127, Trieste, Italy**Italy
JOURNAL: European Journal of Immunology 32 (10): p2773-2782 October, 2002
2002
MEDIUM: print
ISSN: 0014-2980
DOCUMENT TYPE: Article
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RECORD TYPE: Abstract LANGUAGE: English (Item 1 from file: 73) 4/3/2 DIALOG(R)File 73:EMBASE (c) 2010 Elsevier B.V. All rts. reserv. 0081757282 EMBASE/Medline No: 2007191236 Selective therapeutic control of C5a and the terminal complement complex by anti-C5 single-chain Fv in an experimental model of antigen-induced arthritis in rats Fischetti F.; Durigutto P.; Macor P.; ***Marzari R.*** ; Carretta R.; Tedesco F. University of Trieste, Trieste, Italy AUTHOR EMAIL: tedesco@units.it CORRESP. AUTHOR/AFFIL: Tedesco F.: Department of Physiology and Pathology, University of Trieste, via Valerio 28, 34127 Trieste, Italy CORRESP. AUTHOR EMAIL: tedesco@units.it Arthritis and Rheumatism (Arthritis Rheum.) (United States) April 1, 2007, 56/4 (1187-1197) CODEN: ARHEA ISSN: 0004-3591 DOI: 10.1002/art.22492 DOCUMENT TYPE: Journal; Article RECORD TYPE: Abstract LANGUAGE: English SUMMARY LANGUAGE: English NUMBER OF REFERENCES: 56 ? t s4/7/all(Item 1 from file: 5) 4/7/1 DIALOG(R) File 5: Biosis Previews(R) (c) 2010 The Thomson Corporation. All rts. reserv. BIOSIS NO.: 200200576587 16983076 The cleavage site of C5 from man and animals as a common target for neutralizing human monoclonal antibodies: In vitro and in vivo studies AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti Fabio; Gennaro Renato; Marks James D; Bradbury Andrew; Tedesco Francesco (Reprint) AUTHOR ADDRESS: Dipartimento di Fisiologia e Patologia, Universita di Trieste, Via Fleming 22, I-34127, Trieste, Italy**Italy JOURNAL: European Journal of Immunology 32 (10): p2773-2782 October, 2002 2002 MEDIUM: print ISSN: 0014-2980 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English ABSTRACT: The isolation of an anti-C5 single-chain fragment variable (scFv) antibody, TS-A12/22, from a human phage display library, is described. This ***antibody*** inhibits the activation of C5 and the assembly of the terminal complement complex implicated in cell and tissue damage. Using ***antibody*** -sensitized sheep erythrocytes and rabbit red cells as target cells in hemolytic assays, we found that TS-A12/22 inhibited the activation of C5 by the convertases of both classical and alternative pathways. Western blot analysis and competition experiments with synthetic peptides showed that TS-A12/22 reacted with the a chain of C5 and recognized the cleavage site of this complement component by the C5 convertase. As a result, the ***antibody***

splitting of C5 and inhibited the generation of C5a and of the

terminal complement complex. The identification of the ***TS*** - ***A12*** /22 recognition site as a conserved sequence in man, mouse, rat and rabbit enabled the demonstration of in vitro inhibition of complement activity in these species. The scFv TS-A12/22 was tested in a rat model of antigen-induced arthritis and proved to be effective in preventing influx of polymorphonuclear cells into the knee joint and C9 deposition on synovial tissue.

4/7/2 (Item 1 from file: 73) DIALOG(R) File 73: EMBASE (c) 2010 Elsevier B.V. All rts. reserv. EMBASE/Medline No: 2007191236 Selective therapeutic control of C5a and the terminal complement complex by anti-C5 single-chain Fv in an experimental model of antigen-induced arthritis in rats Fischetti F.; Durigutto P.; Macor P.; ***Marzari R.*** ; Carretta R.; Tedesco F. University of Trieste, Trieste, Italy AUTHOR EMAIL: tedesco@units.it CORRESP. AUTHOR/AFFIL: Tedesco F.: Department of Physiology and Pathology, University of Trieste, via Valerio 28, 34127 Trieste, Italy CORRESP. AUTHOR EMAIL: tedesco@units.it Arthritis and Rheumatism (Arthritis Rheum.) (United States) April 1, 2007, 56/4 (1187-1197) CODEN: ARHEA ISSN: 0004-3591 DOI: 10.1002/art.22492 DOCUMENT TYPE: Journal; Article RECORD TYPE: Abstract LANGUAGE: English SUMMARY LANGUAGE: English NUMBER OF REFERENCES: 56

Objective. To determine the role of the terminal complement complex (TCC) in the development of experimental antigen-induced arthritis (AIA) and the therapeutic effects of human anti-C5 single-chain Fv (scFv). Methods. Two different anti-C5 scFv, one that inhibits both release of C5a and assembly of the TCC (TS-A 12/22) and another that selectively blocks formation of the TCC (TS-A 8), were injected at the onset of AIA. The effects of these scFv on disease severity were evaluated for up to 21 days and compared with the effects of injection of an unrelated scFv. AIA was also established in C6-deficient and C6-sufficient PVG rats to obtain further information on the role of the TCC in this model. Results. TS-A 12/22 and TS-A 8 proved to be equally effective in reducing joint swelling, cell counts and tumor necrosis factor alpha levels in synovial lavage fluids, and the degree of histomorphologic changes compared with the effects of the unrelated scFv. TS-A 12/22 and TS-A 8 prevented the deposition of C9 but not that of C3, confirming the ability of the 2 scFv to neutralize C5. Administration of the 2 anti-C5 scFv after AIA onset also reduced disease severity. In C6-deficient rats with AIA, disease activity was reduced markedly compared with that in C6-sufficient rats. Conclusion. These 2 human anti-C5 scFv could represent potential therapeutic reagents to be used in patients with rheumatoid arthritis. In addition, the finding that TS-A 8 was as effective as TS-A 12/22 in reducing disease severity suggests that the TCC is mainly responsible for the joint inflammation and damage observed in AIA. (c) 2007, American College of Rheumatology.

? s (antibod? or immunoglobulin? or hybridoma?)(20n)(c5a or ts(w)a12?)

2724863 ANTIBOD? 1064219 IMMUNOGLOBULIN?

61953 HYBRIDOMA?

14523 C5A

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            5903 A12?
               4 TS(W)A12?
      S_5
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               0 A12?)
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? s (antibod? or immunoglobulin? or hybridoma?)(20n)(c5a) and (ts(w)a12?)
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 8/3/1
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DIALOG(R)File 5:Biosis Previews(R)
(c) 2010 The Thomson Corporation. All rts. reserv.
          BIOSIS NO.: 200200576587
16983076
The cleavage site of C5 from man and animals as a common target for
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AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti Fabio;
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MEDIUM: print
ISSN: 0014-2980
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
 8/3/2
           (Item 1 from file: 73)
DIALOG(R) File 73: EMBASE
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0082118885
              EMBASE/Medline No: 2007532089
  Rheumatoid arthritis and the complement system
  Okroj M.; Heinegard D.; Holmdahl R.; Blom A.M.
 Lund University, Department of Laboratory Medicine, University Hospital
 Malmo, Malmo, Sweden
  AUTHOR EMAIL: Anna.Blom@med.lu.se
  CORRESP. AUTHOR/AFFIL: Blom A.M.: Lund University, Department of
Laboratory Medicine, University Hospital Malmo, Entrance 46, S-205 02 Malmo
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Sweden
 CORRESP. AUTHOR EMAIL: Anna.Blom@med.lu.se
 Annals of Medicine (Ann. Med.) (Norway) November 15, 2007, 39/7
  (517 - 530)
  CODEN: ANMDE
                ISSN: 0785-3890 eISSN: 1651-2219
  PUBLISHER ITEM IDENTIFIER: 780337113
  DOI: 10.1080/07853890701477546
  DOCUMENT TYPE: Journal; Review RECORD TYPE: Abstract
                     SUMMARY LANGUAGE: English
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 NUMBER OF REFERENCES: 139
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         1064219 IMMUNOGLOBULIN?
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DIALOG(R) File 5: Biosis Previews(R)
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